Amendments to the Claims

Claims 1-24 (Cancelled).

- 25. (Currently amended) A composition for delivery of superoxide dismutase (SOD) to neuronal cells, comprising SOD,; linked by a cleavable linker to a neuronal cell targeting component, wherein said neuronal cell targeting component comprises a first domain that binds to a neuronal cell and a second domain that translocates the SOD of the composition into the neuronal cell, and wherein said cleavable linker is selected from the group consisting of (a) a disulfide bridge, and (b) a site for a protease found in neuronal cells.
 - 26. (Cancelled).
 - 27. (Cancelled).
 - 28. (Cancelled).
- 29. (Previously presented). The composition of claim 25 wherein the SOD is bacterial SOD or is derived therefrom.
- 30. (Currently amended). The composition of claim 25 wherein the first domain is selected from the group consisting of
 - (a) neuronal cell binding domains of clostridial toxins; and
 - (b) fragments, variants and derivatives of the domains in (a) that substantially retain the neuronal cell binding activity of the domains of (a).
- 31. (Currently amended) The composition of claim 30 wherein the second domain is selected from the group consisting of

- (a) domains of clostridial neurotoxins that translocate polypeptide sequences into cells, and
- (b) fragments, variants and derivatives of the domains of (a) that substantially retain the translocating activity of the domains of (a).
- 32. (Previously presented) The composition of claim 25 wherein the linker is a disulphide bridge.
- 33. (Currently amended) A pharmaceutical composition for treatment of oxidative damage to neuronal cells comprising the composition of claim 25 and a pharmaceutically acceptable carrier.
 - 34. (Cancelled).
 - 35. (Cancelled).
- 36. (Currently amended) A composition for delivery of an agent a therapeutic agent to neuronal cells, comprising the therapeutic agent; linked by a cleavable linker to a neuronal cell targeting component, wherein said neuronal cell targeting component comprises a first domain that binds to a neuronal cell and a second domain that translocates the therapeutic agent of the composition into the neuronal cell, and wherein said cleavable linker is a disulfide bridge.
 - 37. (Cancelled).
 - 38. (Cancelled).
 - 39. (Cancelled).
 - 40. (Cancelled).
 - 41. (Cancelled).

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- 42. (Previously presented) The composition of claim 25 wherein the cleavable linker is a disulphide bridge between first and second cysteine residues, wherein said first cysteine residue is on the SOD and said second cysteine residue is on the neuronal cell targeting component.
- 43. (Previously presented) The composition of claim 25 wherein the cleavable linker is a site for a protease found in neuronal cells.

This listing of claims will replace all prior versions, and listings of claims in the application.